

## Intended Use

For the quantitative determination of Uric Acid in serum using the Yumizen C230 and Yumizen C240 analyzers. For in vitro diagnostic use only.

### Rx Only.

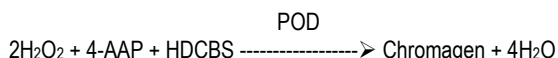
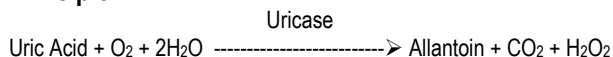
## Clinical Significance

The determination of uric acid in serum is most commonly performed for the diagnosis of gout. Increased uric acid levels are also found in leukemia, polycythemia, familial idiopathic hyperuricemia, and conditions associated with decreased renal function.

## Test Summary

Uric Acid has been determined by phosphotungstate methods,<sup>1</sup> variations of the phosphotungstate method<sup>2</sup> and iron reduction methods.<sup>3,4</sup> The above methodologies are influenced by many substances in their procedures as well as many contaminating substances on glassware, etc.<sup>5</sup> The enzyme Uricase has been widely used for Uric Acid determinations because of its improved specificity.<sup>6,7</sup> Recently, hydrogen peroxide, a by-product of the Uricase-Uric Acid reaction, has been coupled to other enzymatic reactions to yield a colorimetric end product. The present procedure uses the coupling of 4-aminoantipyrine (4-AAP), 2-Hydroxy-3,5-Dichloro-benzenesulfonate (HDCBS), and hydrogen peroxide in the presence of peroxidase to yield a chromagen measured at 520nm.

## Principle



Uric Acid is oxidized by Uricase to allantoin and hydrogen peroxide. HDCBS + 4-AAP + hydrogen peroxide, in the presence of peroxidase, produces a red chromagen that is measured at 520nm. The absorbance at 520nm is proportional to the concentration of Uric Acid in the sample.

## Reagent Composition

Uric Acid reagent: 4-AAP >0.2mM, HDCBS 2mM, Uricase (Microbial) >150 U/L, Peroxidase (horseradish) >2,500 U/L, Buffer, pH 8.1 ± 0.1, Non-reactive stabilizers.

## Reagent Preparation

The reagent is ready to use.

## Reagent Storage and Stability

The reagent set is stored at 2-8°C. Under proper storage the reagent will remain stable until the indicated expiration date.

## Precautions

1. This reagent set is for in vitro diagnostic use only.
2. The reagent should not be used if: The reagent is turbid or contains obvious microbial growth. The reagent blank has an absorbance of 0.500 or greater at 520nm. A pink color is normal for this reagent.
3. All specimens and controls should be handled as potentially infectious, using safe laboratory procedures. (NCCLS M29-T2)<sup>8</sup>

## Specimen Collection and Storage

1. Unhemolyzed serum is recommended.
2. Uric Acid in serum is stable for three days at 2-8°C and up to six months when frozen.<sup>9</sup>
3. Collect specimens per NCCLS document H4-A3.<sup>10</sup>

## Interferences

1. Elevated ascorbic acid levels can result in falsely depressed uric acid values.

2. Lipemic samples may cause falsely elevated uric acid levels.
3. Hemoglobin to 100 mg/dl has been demonstrated to have a negligible effect (<5%) on uric acid values. Hemoglobin greater than 100 mg/dl may cause falsely elevated uric acid values.
4. Bilirubin to 30 mg/dl has been demonstrated to have a negligible effect (<5%) on uric acid results using this method.
5. See Young, et al<sup>11</sup> for other interfering substances.

## Materials Provided

Uric Acid Reagent

## Materials Required but not Provided

1. Yumizen C230 / Yumizen C240 Analyzer
2. Yumizen C230 / Yumizen C240 Operation manual
3. Pointe Chemistry Calibrator, catalog number C7506-50
4. Pointe Chemistry control, catalog number C7592-100

## Test Parameters

Test:	URIC	Chemistry:	Uric Acid
Chemistry No.:	231	Print Name:	Uric Acid
Reaction Type:	Endpoint	Reaction Direction:	Positive
Pri. Wave:	510 nm	Sec. Wave:	670 nm
Decimal.:	0.1	Samp. Type:	Serum
Blank Time:		Reaction Time:	35 37
Unit:	mg/dL	Incubation Time:	0

	Sample Vol.	Aspirated	Diluent	Reagent Vol.	Diluent
Standard;	4	uL	uL	180	uL uL
Decreased;		uL	uL	uL	
Increased;		uL	uL	uL	

Linearity Range (Standard):	0-20	Linearity Limit:	
Linearity Range (Decreased):		Substrate Depletion:	
Linearity Range (Increased):		Mixed Blank Abs.:	-40000 40000
R1 Blank Abs.:	-40000 40000	On-board Stability:	30 Day (s)
Blank Response	-40000 40000	Reagent Alarm Limit:	5
Twin Chemistry:			

Prozone Check:		
Q1:	Q2:	Q3:
Q4:	PC:	ABS:

Use Qualitative Result:	
Range:	Flag:

Slope Offset:			
	Slope	Offset	Unit
	1	0	mg/dL

Pretreatment:	
Pretreat Sample Vol.:	uL Pretreat Reagent Vol.: uL

Ref. Range:			
Sample Type:	Gender:	Age Range:	Ref. Range: Critical Range: Unit:

# Pointe Uric Acid Reagent Set (UV)

## Calibration Setup Parameters

Chem: UA																	
Calibration Setting	<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th>Calibrator</th> <th>Conc.</th> <th>Pos</th> <th>Lot No.</th> </tr> <tr> <td>Water</td> <td>0.0</td> <td>W</td> <td></td> </tr> <tr> <td>Chem Cal</td> <td>*</td> <td>*</td> <td></td> </tr> <tr> <td> </td> <td> </td> <td> </td> <td> </td> </tr> </table>	Calibrator	Conc.	Pos	Lot No.	Water	0.0	W		Chem Cal	*	*					
Calibrator	Conc.	Pos	Lot No.														
Water	0.0	W															
Chem Cal	*	*															
Math Model: Two-Point Linear																	
Factor: Replicates: 2																	
Acceptance Limits																	
Cal Time: 336 hr.																	
Slope Diff: SD:																	
Sensitivity: Repeatability: * User Defined																	
Deter Coeff:																	
Auto Calib.																	
<input type="checkbox"/> Cal Time																	

## Limitations

- If the spectrophotometer being used requires a final volume greater than 1.0ml for accurate reading, use 0.075ml (75ul) of sample to 3.0ml of reagent. Perform the test as described above.
- The procedure described is linear to 20 mg/dl. Samples with values exceeding 20 mg/dl should be diluted 1:1 with saline, re-assayed, and the results multiplied by two.
- Lipemic samples will give falsely elevated results and a serum blank must be run. Serum Blank: Add 0.025ml (25ul) of sample to 1.0ml water. Zero spectrophotometer with water. Read and record absorbance and subtract reading from test absorbance. Calculate as usual.

## Calibration

Use an NIST-traceable serum calibrator. The procedure should be calibrated according to the instrument manufacturer's calibration instructions. If control results are found to be out of range, the procedure should be re-calibrated.

## Calculations (Example)

A = Absorbance

$$\frac{A(\text{Unk})}{A(\text{Std})} \times \text{Conc. of Std (mg/dl)} = \text{Uric Acid (mg/dl)}$$

Example: A (Unk) = 0.126, A (Std) = 0.100, Conc. of Std = 5 mg/dl.

$$\text{Then: } \frac{0.126}{0.100} \times 5 = 6.3 \text{ mg/dl}$$

## SI Units (mM/L)

To convert to mM/L, multiply the result (mg/dl) by 10 to convert dl to L and divide by 168 (the molecular weight of Uric Acid).

$$\text{Mg/dl} \times \frac{10}{168} = \text{mM/L} \quad \text{mg/dl} \times .0595 = \text{mM/L}$$

Example: 6.3mg/dl x .0595 = 0.374mM/L

## Quality Control

Serum controls with known normal and abnormal uric acid values should be run routinely to monitor the validity of the reaction. These controls should be run at least with every working shift in which uric acid determinations are performed. It is strongly recommended that each laboratory establish its own frequency of control determination. Quality control requirements should be performed in conformance with local, state, and/or Federal regulations or accreditation requirements.

## Expected Values

2.5 - 7.7mg/dl<sup>9</sup>

It is strongly recommended that each laboratory establish its own normal range.

## Performance

- Assay Range: 0 - 20 mg/dl
- Comparison: A study was performed between the Yumizen 200 series analyzers and a similar analyzer and method, resulting in a correlation coefficient of 0.998 and the regression equation was  $y=1.015x+0.02$ .
- Precision: Precision studies were performed using the Yumizen 200 series analyzers following a modification of the guidelines which are contained in NCCLS document EP5-T2.<sup>12</sup>

Within Day (n=20)			Day to Day (n=20)		
Mean	S.D.	C.V.%	Mean	S.D.	C.V.%
6.63	0.31	4.6	7.11	0.14	2.0
9.38	0.16	1.7	10.13	0.20	2.0

- Sensitivity: The sensitivity of this reagent was investigated by reading the change in absorbance at 520nm for a saline sample, and two serum samples with known concentrations. Ten replicates of each sample were performed. The results of this investigation indicated that, on the analyzer used, the Uric Acid (Liquid) reagent showed little or no reagent drift on a zero sample. Also, that an absorbance change of 0.015 was approximately equivalent to 1 mg/dl of Uric Acid.

## References

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- Praetorius, E., Poulson, H., Scand. J. Clin. Invest 5:273 (1953).
- NCCLS document "Protection of Laboratory Workers from Infectious Disease Transmitted by Blood, Body Fluids, and Tissue", 2<sup>nd</sup> Ed. (1991).
- Henry, R.J., Clinical Chemistry: Principles and Technics, 2<sup>nd</sup> Ed., Hagerstown (MD), Harper & Row, pp. 531 & 541 (1974).
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- Young, D.S., et al. Clin. Chem. 21:1D (1975).
- NCCLS document "Evaluation of Precision Performance of Clinical Chemistry Devices", 2<sup>nd</sup> Ed. (1992).

## Symbol Key

Use by (YYYY-MM-DD)	<b>LOT</b> Lot and batch code
<b>REF</b> Catalog number	Manufacturer
<b>IVD</b> In vitro diagnostic medical device	Temperature limitation
Consult instructions for use	<b>Rx Only:</b> Prescription Use Only
CE mark	<b>EC REP</b> Authorized representative in the European Community

12-U7581-120

Manufactured by  
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## Certified to Perform Reagents

The Pointe reagents are certified to be manufactured according to specified parameters. Any Pointe reagent product not meeting specifications through its listed expiration date will be remedied immediately without charge.